



THE UNIVERSITY *of* EDINBURGH

## Edinburgh Research Explorer

### **Effects of common genetic variants associated with colorectal cancer risk on survival outcomes after diagnosis: a large population-based cohort study.**

**Citation for published version:**

He, Y, Theodoratou, E, Li, X, Din, FVN, Vaughan-Shaw, P, Svinti, V, Farrington, SM, Campbell, H, Dunlop, MG & Timofeeva, M 2019, 'Effects of common genetic variants associated with colorectal cancer risk on survival outcomes after diagnosis: a large population-based cohort study.', *International Journal of Cancer*.  
<https://doi.org/10.1002/ijc.32550>

**Digital Object Identifier (DOI):**

[10.1002/ijc.32550](https://doi.org/10.1002/ijc.32550)

**Link:**

[Link to publication record in Edinburgh Research Explorer](#)

**Document Version:**

Publisher's PDF, also known as Version of record

**Published In:**

International Journal of Cancer

**General rights**

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.




**Take down policy**

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact [openaccess@ed.ac.uk](mailto:openaccess@ed.ac.uk) providing details, and we will remove access to the work immediately and investigate your claim.



## Short Report

# Effects of common genetic variants associated with colorectal cancer risk on survival outcomes after diagnosis: A large population-based cohort study

Yazhou He <sup>1,2,3</sup>, Evropi Theodoratou<sup>1,3</sup>, Xue Li <sup>3</sup>, Farhat V.N. Din<sup>1,2</sup>, Peter Vaughan-Shaw<sup>2</sup>, Victoria Svinti<sup>2</sup>, Susan M. Farrington<sup>1,2</sup>, Harry Campbell<sup>3</sup>, Malcolm G. Dunlop<sup>1,2</sup> and Maria Timofeeva <sup>1,2</sup>

<sup>1</sup>Cancer Research UK Edinburgh Centre, Medical Research Council Institute of Genetics & Molecular Medicine, Western General Hospital, The University of Edinburgh, Edinburgh, United Kingdom

<sup>2</sup>Colon Cancer Genetics Group, Medical Research Council Human Genetics Unit, Medical Research Council Institute of Genetics & Molecular Medicine, Western General Hospital, The University of Edinburgh, Edinburgh, United Kingdom

<sup>3</sup>Centre for Global Health Research, Usher Institute of Population Health Sciences and Informatics, The University of Edinburgh, Edinburgh, United Kingdom

Genome-wide association studies have thus far identified 130 genetic variants linked to colorectal cancer (CRC) risk ( $r^2 < 0.2$ ). Given their implication in disease causation, and thus plausible biologically effects on cancer-relevant biological pathways, we investigated whether these variants are associated with CRC prognosis and also whether they might provide predictive value for survival outcome. We conducted the analysis in a well-characterized population-based study of 5,675 patients after CRC diagnosis in Scotland. None of the genetic risk variants were associated with either overall survival (OS) or CRC-specific survival. Next, we combined the variants in a polygenic risk score, but again we observed no association between survival outcome and overall genetic susceptibility to CRC risk—as defined by common genetic variants (OS: hazard ratio = 1.00, 95% confidence interval = 0.96–1.05). Furthermore, we found no incremental increase in the discriminative performance when adding these genetic variants to the baseline CRC-survival predictive model of age, sex and stage at diagnosis. Given that our study is well-powered ( $>0.88$ ) to detect effects on survival for 74% of the variants, we conclude that effects of common variants associated with CRC risk which have been identified to date are unlikely to have clinically relevant effect on survival outcomes for patients diagnosed with CRC.

## Introduction

Globally, colorectal cancer (CRC) is the second leading cause of cancer-related deaths, accounting for 9.2% of all cancer-related deaths (0.8 million CRC deaths in 2018).<sup>1</sup> The strongest known

predictor of CRC outcome is stage, but even within one stage, there is considerable heterogeneity in survival. Identification of biomarkers of cancer prognosis can inform clinical management and treatment of disease. Evidence of the family concordance for CRC-

**Additional Supporting Information** may be found in the online version of this article.

**Key words:** common genetic variants, colorectal cancer, survival, cohort study

**Abbreviations:** AJCC: American Joint Committee on Cancer; CI: confidence interval; CRC: colorectal cancer; FDR: false positive rates; HR: hazard ratio; LASSO: least absolute shrinkage and selection operator; MAF: minor allele frequency; PRS: polygenic risk score; SCR: Scottish Cancer Registry; SOCCS: Study of Colorectal Cancer in Scotland

**Conflict of interest:** All authors declare no conflicts of interest.

**Grant sponsor:** MRC Clinical Research Training Fellowship; **Grant number:** MR/M004007/1; **Grant sponsor:** CRUK Career Development Fellowship; **Grant number:** C31250/A22804; **Grant sponsor:** MRC Human Genetics Unit Centre; **Grant numbers:** U127527198, U127527202; **Grant sponsor:** Edinburgh CRUK Cancer Research Centre; **Grant sponsor:** CRUK Program; **Grant number:** C348/A18927

This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

**DOI:** 10.1002/ijc.32550

**History:** Received 3 May 2019; Accepted 17 Jun 2019; Online 4 Jul 2019

**Correspondence to:** Malcolm G. Dunlop, Institute of Genetics and Molecular Medicine, The University of Edinburgh, Western General Hospital, Edinburgh EH4 2XU, United Kingdom, Tel./Fax: +44-131-651-8631, E-mail: malcolm.dunlop@igmm.ed.ac.uk; or Maria Timofeeva, Institute of Genetics and Molecular Medicine, The University of Edinburgh, Western General Hospital, Edinburgh EH4 2XU, United Kingdom, Tel./Fax: +44-131-651-8687, E-mail: maria.timofeeva@igmm.ed.ac.uk

**What's new?**

Genetic variants associated with colorectal cancer (CRC) risk potentially also exert effects on disease prognosis, though little is known about the heritability of CRC survival. In this population-based study, genetic variants of CRC susceptibility were investigated for their relevance in CRC survival based on analyses of variants identified in genome-wide association studies (GWASs). Nonsignificant associations were detected between small numbers of genetic variants and overall survival and CRC-specific survival. Overall, the findings do not support the existence of prognostic effects of common CRC risk variants. Rather, CRC survival may have distinct genetic determinants, warranting separate investigation by GWAS.

specific survival,<sup>2</sup> together with some suggestions of improved survival in cancer patients with a family history compared to patients without a family history,<sup>3</sup> indicates that genetic signature can affect prognosis of CRC patients after diagnosis. Indeed, improved survival for Lynch syndrome patients with germline rare variations in DNA mismatch repair genes is well documented,<sup>4</sup> suggesting that genetic variants associated with CRC pathogenesis may subsequently affect tumor progression. However, very few studies with sufficient power tested roles of common genetic risk variants in CRC prognosis. Previously published smaller studies examined up to 30 CRC risk genetic loci and detected no or little evidence of associations with survival.<sup>5–7</sup> Two recent large meta-analyses of genome-wide association studies (GWAS)<sup>8,9</sup> have identified more than 70 new genetic variants associated with CRC risk. In this analysis, we investigated the association between all previously and newly GWAS-identified common genetic variants and CRC survival.

**Materials and Methods**

We included 5,675 CRC cases (detailed patient selection in Supporting Information Fig. S1) with genome-wide genotyping data and data on age at diagnosis, sex and American Joint Committee on Cancer (AJCC) stage information from a population-based case-control study (Study of Colorectal Cancer in Scotland, SOCCS; 1999-current).<sup>10</sup> Ethics approval was obtained from the MultiCentre Research Ethics committee for Scotland (approval number MREC/01/0/5) and other committees (presented elsewhere<sup>10</sup>). A total of 130 genetic variants identified by previous GWAS studies<sup>8,9</sup> were genotyped or imputed (25/130 variants were directly genotyped). For correlated variants (linkage disequilibrium  $r^2 > 0.2$ ), we selected ones with smaller  $p$  values in the association with CRC risk. Genotyping was conducted using the Illumina HumanHap300, HumanHap240S and OmniExpressExome BeadChip 8v1 arrays. Standard quality control measures were applied as described previously.<sup>9,11,12</sup> Untyped variants were imputed using SHAPEIT v2<sup>13</sup> and IMPUTEv2<sup>14</sup> softwares based on a merged reference panel comprising of 1,000 Genomes Project (phase 1, December 2013 release) and UK10K (April 2014 release) samples. We excluded poorly imputed variants (information measure  $< 0.80$ ) and rare variants (minor allele frequency [MAF]  $< 0.05\%$ ) as presented in the previous publication.<sup>9</sup> Death registration and cause of death was ascertained from the Scottish Cancer Registry (SCR), and patients were prospectively followed up until death or July 1, 2017 (censored date), whichever came first. The survival outcomes included overall survival (OS) and CRC specific survival (CSS). The criteria of assigning cause of death can be found elsewhere.<sup>15</sup> In order to measure

the overall genetic CRC susceptibility, we created polygenic risk scores (PRS) including all the 130 variants based on the number of CRC risk alleles carried by each patient. We employed Cox proportional hazards models to investigate effects of individual variant (additive model) and the PRS on survival outcomes adjusting for age at diagnosis, gender and AJCC stage. Bonferroni correction for multiple testing was adopted and  $p < 0.0005$  was considered statistical significance. We also used the false positive rates (FDR) approach ( $p < 0.05$  was the significance threshold after correction) as a sensitivity analysis.<sup>16</sup> A summary of the 130 included genetic variants is presented in Supporting Information Table S1. Using the method provided by Owzar *et al.*,<sup>17</sup> we calculated the power of variants with various minor allele frequency (MAF) on a range of effects. Stratified analyses were also performed by sex, stage and tumor site. To further explore the potential predictive value of these variants, we applied a least absolute shrinkage and selection operator (LASSO) regression model with 10-fold cross-validation in 70% randomly selected patients (training set) to select predictors.<sup>18</sup> Harrell's concordance indices (C statistic) were calculated to measure the discriminative ability of selected predictors in the remaining 30% patients (test set) and a U-statistic test was adopted to determine the probability that added genetic variants could increase the model concordance.<sup>19</sup>

We also compared hazard ratio (HR) for OS and CSS with the risk results from a recently published meta-analysis of genome-wide association studies of CRC risk.<sup>9</sup> We excluded ambiguous AT and CG variants ( $n = 6$ , rs10161980, rs2186607, rs2696839, rs2732875, rs61336918, rs7398375) from the analysis to avoid bias due to strand differences between studies. For the variants included in the analysis ( $n = 124$ ), we aligned alleles and effect estimates between GWAS on risk and SOCCS survival analysis. We hypothesized that in case of no effects, the CRC risk variants will cause improvement or impairment of OS and CSS in equal proportion (62 variants with effects on risk and survival going in the same direction *vs.* 62 variants with effects on risk and survival going in opposite directions). We further counted all instances of results with similar directions of effects such a HR and OR above 1 or below 1 and compared it to the expected distribution using exact binomial test. Proportions of risk variants associated with worse OS and CSS and corresponding 95% confidence intervals (CIs) were calculated.

All genetic variants were annotated using (i) association with the *cis* gene expression in colon transverse tissue ( $n = 246$ ) from the Genotype-Tissue Expression (GTEx) database,<sup>20</sup> (ii) presence of known and predicted regulatory elements in RegulomeDB database<sup>21</sup> and (iii) predicted effect on the structure and function of a protein as implemented in SIFT<sup>22</sup> and PolyPhen-2.<sup>23</sup>

**Table 1.** Basic characteristics of included CRC cases

Characteristics	SOCCS CRC cases (n = 5,675)
Age at diagnosis (years) <sup>1</sup>	64.5 (54.6–71.6)
Sex	
Male	3,235 (57.0%)
Female	2,440 (43.0%)
AJCC stage	
I	1,005 (17.7%)
II	1,891 (33.3%)
III	1,995 (35.2%)
IV	784 (13.8%)
Site	
Colon	3,392 (59.8%)
Rectum	2,201 (38.8%)
Colon and rectum	16 (0.3%)
Unknown	66 (1.2%)
Follow-up time (years) <sup>1</sup>	5.09 (2.43–11.42)
No. of all-cause deaths	1,918 (33.8%)
No. of CRC-related deaths	1,358 (23.9%)

<sup>1</sup>Median and quartiles in parenthesis.

Abbreviations: AJCC, American Joint Committee on Cancer; CRC, colorectal cancer; SOCCS, Study of Colorectal Cancer in Scotland.

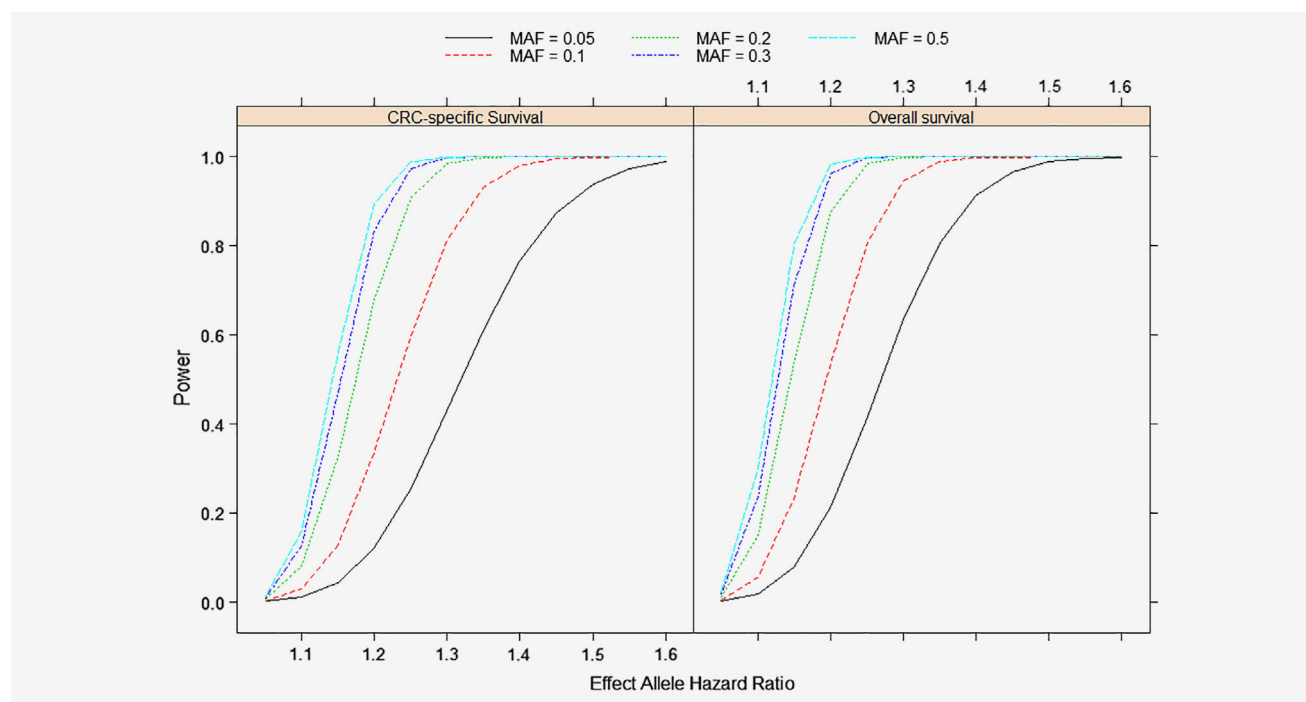
### Data availability

The data that support the findings of our study are available upon reasonable request from the corresponding authors. The data are not publicly available due to privacy or ethical restrictions.

### Results and Discussion

The basic characteristics of included CRC patients are summarized in Table 1. In total, 1,918 patients (34%) died during follow-up. With 5,675 CRC cases, our study had 88% power to detect a hazard ratio of 1.20 on OS for 97/130 (74%) variants (MAF > 0.15) at the significance level of 0.0005 (a power curve for other effects is shown in Fig. 1). Overall, we observed 14 genetic variants associated with OS and 10 with CSS at nominal statistical significance ( $p < 0.05$ ) with six variants (rs10994860, rs12143541, rs3217810, rs34405347, rs6065668, rs847208) being associated with both OS and CSS. However, none of the variants remained statistically significant after Bonferroni or FDR correction. The summary results for variants with nominal significance are presented in Table 2. Stratified analyses by stage, sex and tumor site did not identify any statistically significant associations after multiple-testing correction either (Supporting Information Tables S2–S4). With regard to overall genetic susceptibility to CRC, no statistically significant association was observed between the PRS and OS or CSS (Table 2). The LASSO regression model selected six variants for OS in addition to age at diagnosis, sex and AJCC stage to minimize prediction error in the train set. However, nearly no incremental predictive improvement was observed compared to the model without genetic variants in the test set (C-statistic for OS: 0.73282 vs. 0.73277, U-statistic test:  $p = 0.322$ ). Similar results were found for CSS (Supporting information Table S5).

We used a concept of statistical significance as decision criteria to define if the risk variants have an effect on survival. This concept has been criticized in the literature as subjective and commonly misused.<sup>24,25</sup> We additionally looked into



**Figure 1.** Power curve for overall and CRC-specific survival using additive model. Abbreviations: CRC, colorectal cancer; MAF, minor allele frequency. [Color figure can be viewed at wileyonlinelibrary.com]

Table 2. Summarized results of association between variants at nominal significance ( $p < 0.05$ ) and PRS with CRC survival

Genetic marker				Overall survival			CRC-specific survival		
Variant	Gene	EA	MAF <sup>2</sup>	HR <sup>1</sup> (95% CI)	<i>p</i> value	Pfdr	HR <sup>1</sup> (95% CI)	<i>p</i> value	Pfdr
rs10161980	AL139383.1	G	0.40	1.08 (1.01–1.15)	0.019	0.251	1.06 (0.98–1.14)	0.160	0.832
rs10994860	A1CF	T	0.17	1.11 (1.02–1.20)	0.016	0.251	1.13 (1.02–1.24)	0.018	0.603
rs11196171	TCF7L2	G	0.22	0.91 (0.83–0.98)	0.017	0.251	0.93 (0.85–1.03)	0.161	0.832
rs12143541	TTC22	G	0.15	1.13 (1.04–1.24)	0.006	0.251	1.13 (1.02–1.25)	0.023	0.603
rs16959063	FMN1	A	0.01	0.72 (0.53–0.97)	0.034	0.380	0.83 (0.59–1.16)	0.273	0.832
rs174537	MYRF	T	0.34	1.08 (1.01–1.16)	0.019	0.251	1.05 (0.97–1.14)	0.218	0.832
rs2696839	Intergenic	C	0.48	0.95 (0.90–1.02)	0.143	0.642	0.93 (0.86–1.00)	0.048	0.603
rs3087967	C11orf53	C	0.33	0.92 (0.86–0.99)	0.017	0.251	0.93 (0.86–1.00)	0.062	0.616
rs3217810	CCND2	T	0.14	1.13 (1.02–1.25)	0.016	0.251	1.13 (1.00–1.27)	0.044	0.603
rs3217874	CCND2	T	0.42	1.07 (1.00–1.14)	0.050	0.438	1.06 (0.98–1.14)	0.139	0.823
rs34405347	Intergenic	G	0.08	0.84 (0.74–0.95)	0.007	0.251	0.85 (0.73–0.99)	0.042	0.603
rs35509282	Intergenic	A	0.12	0.94 (0.86–1.04)	0.256	0.772	0.88 (0.78–0.99)	0.033	0.603
rs4759277	LRP1	A	0.38	1.07 (1.00–1.14)	0.051	0.438	1.09 (1.01–1.18)	0.028	0.603
rs6065668	Intergenic	T	0.28	0.90 (0.83–0.96)	0.003	0.251	0.89 (0.82–0.97)	0.011	0.603
rs7495132	CRTC3	T	0.12	1.03 (0.93–1.13)	0.602	0.888	1.13 (1.01–1.26)	0.032	0.603
rs847208	LINC01081	A	0.37	0.93 (0.88–1.00)	0.042	0.416	0.92 (0.85–0.99)	0.027	0.603
rs9537521	AL139383.1	A	0.38	1.10 (1.02–1.17)	0.008	0.251	1.06 (0.98–1.15)	0.153	0.832
rs9929218	CDH1	A	0.28	0.93 (0.86–0.99)	0.035	0.380	0.96 (0.88–1.04)	0.294	0.836
PRS		NA		1.00 (0.95–1.04)	0.864	NA	1.03 (0.97–1.08)	0.340	NA

<sup>1</sup>Hazard ratios are derived from Cox regression models adjusted for age at diagnosis, sex and AJCC stage.

<sup>2</sup>Minor allele frequency is the minor allele prevalence in SOCCS.

Abbreviations: CI, confidence interval; CRC, colorectal cancer; EA, effect alleles; HR, hazard ratio; MAF, minor allele frequency; NA, not available; Pfdr, *p* values after correction for false positive rates; PRS, polygenic risk score.

direction of effects to overcome limitations of statistical significance. We hypothesized that in case of no effects, the CRC risk variants will cause improvement or impairment of OS and CSS in equal proportion. Ambiguous AT and CG genetic variants were excluded and out of 124 tested variants only 52 risk variants were also associated with worse OS (proportion of risk variants associated with worse OS 41%, 95% CI = 33–51%) and 58 were associated with decrease in CSS (proportion of risk variants associated with worse CSS 47%, 95% CI = 38–56%). Though not reaching suggested significance level ( $p \leq 0.05$ ), these results are consistent with directions of effects observed in previous studies.<sup>3,4</sup>

None of the nominally significant genetic variants have known detrimental clinically relevant effects on gene function (Supporting Information Table S6). Rs3087967, which is located 3'UTP of C11orf53, is known to be associated with higher *COLCA2* and *C11orf53* expression in colon transverse tissue for C allele.<sup>9,20</sup> However, little is known about *COLCA2* and C11orf53 functions. rs4759277, which is associated with worse OS and located within intron region of *LRP1*, is likely to affect binding of transcription factors and associated with LRP1 gene expression in tibial artery and sun-exposed skin.<sup>20</sup> Another variant with nominally significant effects on both OS and CSS is rs10994860 variant. It is located 5'UTP of APOBEC1 complementation factor (*A1CF*) and likely to affect binding (regulomeDB score 2b). The same variant has been

previously associated with estimated glomerular filtration rate (eGFR), a measure of the kidney's filtration ability in serum.<sup>26</sup>

This is the first study capturing all currently identified CRC risk genetic loci ( $n = 92$ ) and investigating their associations with survival outcomes in a population-based study. Our results indicate that overall genetic CRC susceptibility measured by GWAS-identified variants is not statistically significantly associated with survival after CRC diagnosis. With regard to each variant, our study, which had acceptable power for  $HR > 1.2$ , found multiple variants (14 for OS and 10 for CSS) associated with survival outcomes at  $p < 0.05$ , although the significance fails to survive correction for multiple testing.

Similarly, previous studies identified some CRC-risk variants<sup>27,28</sup> that might be associated with survival after CRC diagnosis, these findings were not immune to false-positive results from multiple testing. A widely studied variant, rs9929218 lies in the intron of *CDH1* gene encoding E-cadherin, the loss of function of which can cause tumor progression and metastasis.<sup>29</sup> Previous studies reported that the CRC-risk decreasing allele (A) is statistically significantly associated with poor survival outcomes.<sup>27,28</sup> However, in contrast, we observed a potentially favorable effect (though not statistically significant after multiple-testing correction) of the A allele on OS in our study (i.e., the direction of CRC risk and prognosis are consistent in our study). Smith *et al.* reported that the A allele is significantly associated with poor response to chemotherapy, implying a



possible gene  $\times$  therapy interaction for this variant.<sup>28</sup> However, our study is limited by data on response to chemotherapy being unavailable so that this could not be explored further. Notably, there has been other evidence showing that rs9929218 may modify CRC susceptibility by interacting with other factors such as height and alcohol consumption.<sup>30</sup> Investigation of possible gene–environment interactions should be considered in future efforts with large sample sizes to further dissect the prognostic effect of this variant in CRC patients. Consistently, when looking for overall direction of effects among 124 tested variants we noted little evidence of potentially detrimental effects of CRC risk on survival with only 41 and 47% of risk variants showing association with poor (HR > 1) OS and CSS in our study.

Previously we have shown that SOCCS study is representative of British and Scottish populations and cases from SOCCS cluster tightly with population-based controls from SOCCS and Generation Scotland.<sup>31,32</sup> The allele frequencies of studied genetic variants are in range expected for European populations (Supporting Information Table S1). However, the results may be not generalizable to populations of cancer patients where substantial differences in allele frequencies and/or treatment and disease management are anticipated.

In conclusion, our study finds that overall genetic susceptibility to CRC captured by known CRC risk variants is not statistically significantly associated with survival outcomes of

CRC. However, possible roles of each variant in CRC progression remain to be explored. Our study indicated that the heritable variation of patient survival may have distinct genetic determinants from CRC susceptibility. The previous GWAS on CRC survival with 3,494 cases identified no variants at genome-wide significance ( $p < 5E-8$ ),<sup>33</sup> meriting future collaborative efforts of aggregating larger CRC cohorts to illuminate genetic structure of survival outcomes for CRC patients.

## Acknowledgements

We acknowledge the excellent technical support from Stuart Reid and Marion Walker. We are grateful to Donna Markie and all those who continue to contribute to recruitment, data collection, and data curation for the Study of Colorectal Cancer in Scotland studies. We acknowledge that these studies would not be possible without the patients and surgeons who take part. We acknowledge the expert support on sample preparation from the Genetics Core of the Edinburgh Wellcome Trust Clinical Research Facility. We are grateful for the provision of public data from the GTEx consortium. The data used for the analyses described in this article were obtained from the GTEx Portal on 05/27/2019. This work was supported by CRUK Program grant C348/A18927 (MGD). It was also supported by funding for the infrastructure and staffing of the Edinburgh CRUK Cancer Research Centre. This work was also funded by a grant to MGD as Project Leader with the MRC Human Genetics Unit Centre Grant (U127527202 and U127527198 from 1/4/18). ET is supported by a CRUK Career Development Fellowship (C31250/A22804). PVS was supported by MRC Clinical Research Training Fellowship (MR/M004007/1), a research fellowship from the Harold Bridges Bequest and by the Melville Trust for the Care and Cure of Cancer. The work received support from COST Action BM1206.

## References

- Bray F, Ferlay J, Soerjomataram I, et al. Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin* 2018;68: 394–424.
- Lindstrom LS, Hall P, Hartman M, et al. Familial concordance in cancer survival: a Swedish population-based study. *Lancet Oncol* 2007;8: 1001–6.
- Lee M, Reilly M, Lindstrom LS, et al. Differences in survival for patients with familial and sporadic cancer. *Int J Cancer* 2017;140:581–90.
- Watson P, Lin KM, Rodriguez-Bigas MA, et al. Colorectal carcinoma survival among hereditary nonpolyposis colorectal carcinoma family members. *Cancer* 1998;83:259–66.
- Phipps AI, Newcomb PA, Garcia-Albeniz X, et al. Association between colorectal cancer susceptibility loci and survival time after diagnosis with colorectal cancer. *Gastroenterology* 2012;143: 51–4.e4.
- Dai J, Gu J, Huang M, et al. GWAS-identified colorectal cancer susceptibility loci associated with clinical outcomes. *Carcinogenesis* 2012;33: 1327–31.
- Tenesa A, Theodoratou E, Din FV, et al. Ten common genetic variants associated with colorectal cancer risk are not associated with survival after diagnosis. *Clin Cancer Res* 2010;16: 3754–9.
- Huyghe JR, Bien SA, Harrison TA, et al. Discovery of common and rare genetic risk variants for colorectal cancer. *Nat Genet* 2019;51:76–87.
- Law PJ, Timofeeva M, Fernandez-Rozadilla C, et al. Association analyses identify 31 new risk loci for colorectal cancer susceptibility. *Nat Commun* 2019;10:2154.
- Theodoratou E, Kyle J, Cetnarskyj R, et al. Dietary flavonoids and the risk of colorectal cancer. *Cancer Epidemiol Biomarkers Prev* 2007;16:684–93.
- Anderson CA, Pettersson FH, Clarke GM, et al. Data quality control in genetic case-control association studies. *Nat Protoc* 2010;5: 1564–73.
- He Y, Timofeeva M, Farrington SM, et al. Exploring causality in the association between circulating 25-hydroxyvitamin D and colorectal cancer risk: a large Mendelian randomisation study. *BMC Med* 2018;16:142.
- Delaneau O, Marchini J, Zagury JF. A linear complexity phasing method for thousands of genomes. *Nat Methods* 2011;9:179–81.
- Howie BN, Donnelly P, Marchini J. A flexible and accurate genotype imputation method for the next generation of genome-wide association studies. *PLoS Genet* 2009;5:e1000529.
- Theodoratou E, Farrington SM, Timofeeva M, et al. Genome-wide scan of the effect of common nsSNPs on colorectal cancer survival outcome. *Br J Cancer* 2018;119:988–93.
- Benjamini Y, Hochberg Y. Controlling the false discovery rate: a practical and powerful approach to multiple testing. *J R Stat Soc B Methodol* 1995; 57:289–300.
- Owzar K, Li Z, Cox N, et al. Power and sample size calculations for SNP association studies with censored time-to-event outcomes. *Genet Epidemiol* 2012;36:538–48.
- Tibshirani R. Regression shrinkage and selection via the lasso. *J R Stat Soc B Methodol* 1996;58: 267–88.
- Harell F, Williams S. Rank Correlation for Paired Predictors with a Possibly Censored Response, and Integrated Discrimination Index. *Hmisc version 39-3*; 2012: 208–11. [cited January 9, 2019]; Available at: <http://cran.r-project.org/web/packages/Hmisc/Hmisc.pdf>.
- GTEx. [cited 27 May 2019]; Available at: <https://www.gtexportal.org/home/>.
- Boyle AP, Hong EL, Hariharan M, et al. Annotation of functional variation in personal genomes using RegulomeDB. *Genome Res* 2012;22: 1790–7.
- Kumar P, Henikoff S, Ng PC. Predicting the effects of coding non-synonymous variants on protein function using the SIFT algorithm. *Nat Protoc* 2009;4:1073–81.
- Adzhubei IA, Schmidt S, Peshkin L, et al. A method and server for predicting damaging missense mutations. *Nat Methods* 2010;7: 248–9.
- Amrhein V, Greenland S, McShane B. Scientists rise up against statistical significance. *Nature* 2019;567:305–7.
- Wasserstein RL, Lazar NA. The ASA's statement on p-values: context, process, and purpose. *Am Stat* 2016;70:129–33.
- Pattaro C, Teumer A, Gorski M, et al. Genetic associations at 53 loci highlight cell types and

- biological pathways relevant for kidney function. *Nat Commun* 2016;7:10023.
27. Abuli A, Lozano JJ, Rodriguez-Soler M, et al. Genetic susceptibility variants associated with colorectal cancer prognosis. *Carcinogenesis* 2013; 34:2286–91.
  28. Smith CG, Fisher D, Harris R, et al. Analyses of 7,635 patients with colorectal cancer using independent training and validation cohorts show that rs9929218 in CDH1 is a prognostic marker of survival. *Clin Cancer Res* 2015;21:3453–61.
  29. Takeichi M. Cadherin cell adhesion receptors as a morphogenetic regulator. *Science* 1991;251: 1451–5.
  30. Figueiredo JC, Lewinger JP, Song C, et al. Genotype-environment interactions in microsatellite stable/microsatellite instability-low colorectal cancer: results from a genome-wide association study. *Cancer Epidemiol Biomarkers Prev* 2011;20: 758–66.
  31. Timofeeva MN, Kinnersley B, Farrington SM, et al. Recurrent coding sequence variation explains only a small fraction of the genetic architecture of colorectal cancer. *Sci Rep* 2015;5: 16286.
  32. Dunlop MG, Dobbins SE, Farrington SM, et al. Common variation near CDKN1A, POLD3 and SHROOM2 influences colorectal cancer risk. *Nat Genet* 2012;44:770–6.
  33. Phipps AI, Passarelli MN, Chan AT, et al. Common genetic variation and survival after colorectal cancer diagnosis: a genome-wide analysis. *Carcinogenesis* 2016;37:87–95.

# 24 days of stem cells

---

Shape the future of stem cell innovation  
**October 1- November 1, 2019**

**Join us for 24 Days of Stem Cells; a premiere virtual event featuring the latest advances in stem cell research.**

This year's format will feature a new hour of cutting edge content every week day starting October 1st. Attend the sessions that are most relevant to your work - at your convenience and at your pace.

During the 24-day long event, you can:

- Access leading scientific presentations from thought leaders around the world
- Watch live training demonstrations from our stem cell experts
- Download key stem cell tools and resources
- Complete weekly challenges to earn points towards certification and prizes

**Register today at**  
**[www.24daysofstemcells.com](http://www.24daysofstemcells.com)**

**ThermoFisher**  
SCIENTIFIC

**WILEY**